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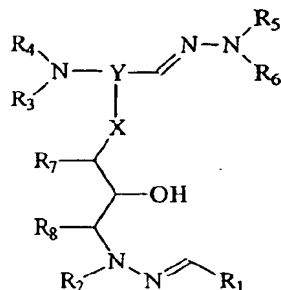
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(54) Organophotoreceptor with charge transport compound with hydrazone groups

(57) This invention relates to a novel organophotoreceptor that comprises at least one photoconductive element comprising

a) a novel charge transport compound having the formula



where R_1 is a carbazole group, a julolidine group, or a p- (N,N-disubstituted) arylamine, R_2 , R_3 , R_4 , R_5 and R_6 are, independently, an alkyl group or an aryl group, R_7 and R_8 are, independently, hydrogen, an alkyl group, or an aryl group, X is oxygen, sulfur, or a NR' group where R' is hydrogen, an alkyl, or an aryl group, and Y is a trivalent aryl group;
(b) a charge generating compound; and

wherein the at least one photoconductive element is carried on an electrically conductive substrate.

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Description

[0001] This application claims priority to copending U.S. Provisional Patent Applications serial number 60/421,182 filed on October 28, 2002 to Getautis et al., entitled "Electrophotographic Photoreceptor With Novel Charge Transport Compound," incorporated herein by reference.

[0002] This invention relates to organophotoreceptors suitable for use in electrophotography and, more specifically, to organophotoreceptors having an improved charge transport compound having at least two hydrazone groups.

[0003] In electrophotography, an organophotoreceptor in the form of a plate, disk, sheet, belt, drum or the like having an electrically insulating photoconductive element on an electrically conductive substrate is imaged by first uniformly electrostatically charging the surface of the photoconductive layer, and then exposing the charged surface to a pattern of light. The light exposure selectively dissipates the charge in the illuminated areas where light strikes the surface, thereby forming a pattern of charged and uncharged areas, referred to as a latent image. A liquid or solid toner is then provided in the vicinity of the latent image, and toner droplets or particles deposit in the vicinity of either the charged or uncharged areas to create a toned image on the surface of the photoconductive layer. The resulting toned image can be transferred to a suitable ultimate or intermediate receiving surface, such as paper, or the photoconductive layer can operate as an ultimate receptor for the image. The imaging process can be repeated many times to complete a single image, for example, by overlaying images of distinct color components or effect shadow images, such as overlaying images of distinct colors to form a full color final image, and/or to reproduce additional images.

[0004] Both single layer and multilayer photoconductive elements have been used. In single layer embodiments, a charge transport material and charge generating material are combined with a polymeric binder and then deposited on the electrically conductive substrate. In multilayer embodiments, the charge transport material and charge generating material are present in the element in separate layers, each of which can optionally be combined with a polymeric binder, deposited on the electrically conductive substrate. Two arrangements are possible. In one two-layer arrangement (the "dual layer" arrangement), the charge generating layer is deposited on the electrically conductive substrate and the charge transport layer is deposited on top of the charge generating layer. In an alternate two-layer arrangement (the "inverted dual layer" arrangement), the order of the charge transport layer and charge generating layer is reversed.

[0005] In both the single and multilayer photoconductive elements, the purpose of the charge generating material is to generate charge carriers (i.e., holes and/or electrons) upon exposure to light. The purpose of the charge transport material is to accept at least one type of these charge carriers, generally holes, and transport them through the charge transport layer in order to facilitate discharge of a surface charge on the photoconductive element. The charge transport material can be a charge transport compound, an electron transport compound, or a combination of both. When a charge transport compound is used, the charge transport compound accepts the hole carriers and transports them through the layer with the charge transport compound. When an electron transport compound is used, the electron transport compound accepts the electron carriers and transports them through the layer with the electron transport compound.

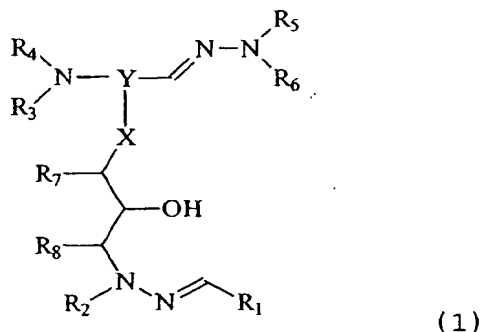
[0006] According to the present invention there is provided an organophotoreceptor, an electrophotographic imaging apparatus, an electrophotographic imaging process, and a charge transport compound, as set forth in the appended claims.

[0007] Preferred features of the invention will be apparent from the dependent claims, and the description which follows.

[0008] This invention provides novel organophotoreceptors having good electrostatic properties such as high V_{acc} and low V_{dis} .

[0009] In a first aspect, an organophotoreceptor comprises at least a photoconductive element comprising:

- a) a charge transport compound having the formula



where R_1 is a carbazole group, a julolidine group, or a p-(N,N-disubstituted)arylamine, R_2 , R_3 , R_4 , R_5 and R_6 are, independently, an alkyl group or an aryl group, R_7 and R_8 are, independently, hydrogen, an alkyl group, or an aryl group, X is oxygen, sulfur, or a NR' group where R' is hydrogen, an alkyl, or an aryl group, and Y is a aryl group; and

(b) a charge generating compound;
wherein the photoconductive element is on an electrically conductive substrate.

[0010] The organophotoreceptor may be provided in the form of a plate, a flexible belt, a flexible disk, a sheet, a rigid drum, or a sheet around a rigid or compliant drum. In one embodiment, the organophotoreceptor includes: (a) a photoconductive element comprising the charge transport compound, the charge generating compound, the electron transport compound, and a polymeric binder; and (b) the electrically conductive substrate.

[0011] In a second aspect, the invention features an electrophotographic imaging apparatus that comprises (a) a light imaging component; and (b) the above-described organophotoreceptor oriented to receive light from the light imaging component. The apparatus can further comprise a liquid toner dispenser. The method of electrophotographic imaging with photoreceptors containing the above noted charge transport compounds is also described.

[0012] In a third aspect, the invention features an electrophotographic imaging process comprising (a) applying an electrical charge to a surface of the above-described organophotoreceptor; (b) imagewise exposing the surface of the organophotoreceptor to radiation to dissipate charge in selected areas and thereby form a pattern of at least relatively charged and uncharged areas on the surface; (c) contacting the surface with a toner, such as a liquid toner that includes a dispersion of colorant particles in an organic liquid, to create a toned image; and (d) transferring the toned image to a substrate.

[0013] In a fourth aspect, the invention features a charge transport compound having the general formula (1) above.

[0014] The invention provides suitable charge transport compounds for organophotoreceptors featuring a combination of good mechanical and electrostatic properties. These photoreceptors can be used successfully with liquid toners to produce high quality images. The high quality of the imaging system can be maintained after repeated cycling.

[0015] Other features and advantages of the invention will be apparent from the following description of the particular embodiments thereof, and from the claims.

[0016] An organophotoreceptor as described herein has an electrically conductive substrate and a photoconductive element comprising a charge generating compound and a charge transport compound having two hydrazone groups linked through an aryl group. These charge transport compounds have desirable properties as evidenced by their performance in organophotoreceptors for electrophotography. In particular, the charge transport compounds of this invention have high charge carrier mobilities and good compatibility with various binder materials, can be cross-linked in both the single and multilayer photoconductive elements, and possess excellent electrophotographic properties. The organophotoreceptors according to this invention generally have a high photosensitivity, a low residual potential, and a high stability with respect to cycle testing, crystallization, and organophotoreceptor bending and stretching. The organophotoreceptors are particularly useful in laser printers and the like as well as photocopiers, scanners and other electronic devices based on electrophotography. The use of these charge transport compounds is described in more detail below in the context of laser printer use, although their application in other devices operating by electrophotography can be generalized from the discussion below.

[0017] To produce high quality images, particularly after multiple cycles, it is desirable for the charge transport compounds to form a homogeneous solution with the polymeric binder and remain approximately homogeneously distributed through the organophotoreceptor material during the cycling of the material. In addition, it is desirable to increase the amount of charge that the charge transport material, such as a charge transport compound, can accept (indicated

by a parameter known as the acceptance voltage or " V_{acc} "), and to reduce retention of that charge upon discharge (indicated by a parameter known as the discharge voltage or " V_{dis} ").

[0018] There are many charge transport compounds available for electrophotography. Examples of charge transport compounds include, for example, pyrazoline derivatives, fluorene derivatives, oxadiazole derivatives, stilbene derivatives, hydrazone derivatives, carbazole hydrazone derivatives, triaryl amines, polyvinyl carbazole, polyvinyl pyrene, polyacenaphthylene, or multi-hydrazone compounds comprising at least two hydrazone groups and at least two groups selected from the group consisting of p-(N,N-disubstituted) arylamine such as triphenylamine and heterocycles such as carbazole, julolidine, phenothiazine, phenazine, phenoxazine, phenoxathin, thiazole, oxazole, isoxazole, dibenzo (1,4)dioxine, thianthrene, imidazole, benzothiazole, benzotriazole, benzoxazole, benzimidazole, quinoline, isoquinoline, quinoxaline, indole, indazole, pyrrole, purine, pyridine, pyridazine, pyrimidine, pyrazine, triazole, oxadiazole, tetrazole, thiadiazole, benzisoxazole, benzisothiazole, dibenzofuran, dibenzothiophene, thiophene, thianaphthene, quinaldine, or cinnoline. However, there is a need for other charge transport compounds to meet the various requirements of particular electrophotography applications.

[0019] In electrophotography applications, a charge generating compound within an organophotoreceptor absorbs light to form electron-hole pairs. These electron-hole pairs can be transported over an appropriate time frame under a large electric field to discharge locally a surface charge that is generating the field. The discharge of the field at a particular location results in a surface charge pattern that essentially matches the pattern drawn with the light. This charge pattern then can be used to guide toner deposition. The charge transport compounds described herein are especially effective at transporting charge, and in particular holes from the electron-hole pairs formed by the charge generating compound. In some embodiments, a specific electron transport compound can also be used along with the charge transport compound.

[0020] The layer or layers of materials containing the charge generating compound and the charge transport compounds are within an organophotoreceptor. To print a two dimensional image using the organophotoreceptor, the organophotoreceptor has a two dimensional surface for forming at least a portion of the image. The imaging process then continues by cycling the organophotoreceptor to complete the formation of the entire image and/or for the processing of subsequent images.

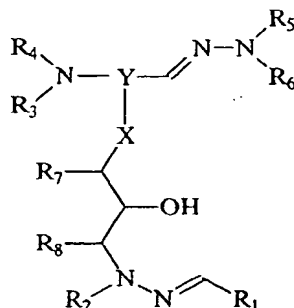
[0021] The organophotoreceptor may be provided in the form of a plate, a flexible belt, a disk, a rigid drum, a sheet around a rigid or compliant drum, or the like. The charge transport compound can be in the same layer as the charge generating compound and/or in a different layer from the charge generating compound. Additional layers can be used also, as described further below.

[0022] In some embodiments, the organophotoreceptor material comprises, for example: (a) a charge transport layer comprising the charge transport compound and a polymeric binder; (b) a charge generating layer comprising the charge generating compound and a polymeric binder; and (c) the electrically conductive substrate. The charge transport layer may be intermediate between the charge generating layer and the electrically conductive substrate. Alternatively, the charge generating layer may be intermediate between the charge transport layer and the electrically conductive substrate. In further embodiments, the organophotoreceptor material has a single layer with both a charge transport compound and a charge generating compound within a polymeric binder.

[0023] The organophotoreceptors can be incorporated into an electrophotographic imaging apparatus, such as laser printers. In these devices, an image is formed from physical embodiments and converted to a light image that is scanned onto the organophotoreceptor to form a surface latent image. The surface latent image can be used to attract toner onto the surface of the organophotoreceptor, in which the toner image is the same or the negative of the light image projected onto the organophotoreceptor. The toner can be a liquid toner or a dry toner. The toner is subsequently transferred, from the surface of the organophotoreceptor, to a receiving surface, such as a sheet of paper. After the transfer of the toner, the entire surface is discharged, and the material is ready to cycle again. The imaging apparatus can further comprise, for example, a plurality of support rollers for transporting a paper receiving medium and/or for movement of the photoreceptor, a light imaging component with suitable optics to form the light image, a light source, such as a laser, a toner source and delivery system and an appropriate control system.

[0024] An electrophotographic imaging process generally can comprise (a) applying an electrical charge to a surface of the above-described organophotoreceptor; (b) imagewise exposing the surface of the organophotoreceptor to radiation to dissipate charge in selected areas and thereby form a pattern of charged and uncharged areas on the surface; (c) exposing the surface with a toner, such as a liquid toner that includes a dispersion of colorant particles in an organic liquid to create a toner image, to attract toner to the charged or discharged regions of the organophotoreceptor; and (d) transferring the toner image to a substrate.

[0025] As described herein, an organophotoreceptor comprises a charge transport compound having the formula



where R_1 is a carbazole group, a julolidine group, or a p-(N,N-disubstituted)arylamine, R_2 , R_3 , R_4 , R_5 and R_6 are, independently, an alkyl group or an aryl group, R_7 and R_8 are, independently, hydrogen, an alkyl group or an aryl group, X is oxygen, sulfur, or a NR' group where R' is hydrogen, an alkyl, or an aryl group, and Y is an aryl group.

[0026] Substitution is liberally allowed on the chemical groups to effect various physical effects on the properties of the compounds, such as mobility, sensitivity, solubility, stability, and the like, as is known generally in the art. In the description of chemical substituents, there are certain practices common to the art that are reflected in the use of language. The term group indicates that the generically recited chemical entity (e.g., alkyl group, phenyl group, julolidine group, (N,N-disubstituted) arylamine group, etc.) may have any substituent thereon which is consistent with the bond structure of that group. For example, where the term 'alkyl group' is used, that term would not only include unsubstituted linear, branched and cyclic alkyls, such as methyl, ethyl, isopropyl, tert-butyl, cyclohexyl, dodecyl and the like, but also substituents such as hydroxyethyl, cyanobutyl, 1,2,3-trichloropropane, and the like. However, as is consistent with such nomenclature, no substitution would be included within the term that would alter the fundamental bond structure of the underlying group. For example, where a phenyl group is recited, substitution such as 1-hydroxyphenyl, 2,4-fluorophenyl, orthocyanophenyl, 1,3,5-trimethoxyphenyl and the like would be acceptable within the terminology, while substitution of 1,1,2,2,3,3-hexamethylphenyl would not be acceptable as that substitution would require the ring bond structure of the phenyl group to be altered to a nonaromatic form because of the substitution. Similarly, when referring to carbazole group or julolidine group, the compound or substituent cited will include any substitution that does not substantively alter the chemical nature of the carbazole ring or the julolidine ring in the formula. When referring p-(N,N-disubstituted)arylamine group, the two substituents attached to the nitrogen may be any group that will not substantively alter the chemical nature of the amine group. Where the term moiety is used, such as alkyl moiety or phenyl moiety, that terminology indicates that the chemical material is not substituted. Where the term alkyl moiety is used, that term represents only an unsubstituted alkyl hydrocarbon group, whether branched, straight chain, or cyclic.

Organophotoreceptors

[0027] The organophotoreceptor may be, for example, in the form of a plate, a sheet, a flexible belt, a disk, a rigid drum, or a sheet around a rigid or compliant drum, with flexible belts and rigid drums generally being used in commercial embodiments. The organophotoreceptor may comprise, for example, an electrically conductive substrate and on the electrically conductive substrate a photoconductive element in the form of one or more layers. The photoconductive element can comprise both a charge transport compound and a charge generating compound in a polymeric binder, which may or may not be in the same layer, as well as an electron transport compound in some embodiments. For example, the charge transport compound and the charge generating compound can be in a single layer. In other embodiments, however, the photoconductive element comprises a bilayer construction featuring a charge generating layer and a separate charge transport layer. The charge generating layer may be located intermediate between the electrically conductive substrate and the charge transport layer. Alternatively, the photoconductive element may have a structure in which the charge transport layer is intermediate between the electrically conductive substrate and the charge generating layer.

[0028] The electrically conductive substrate may be flexible, for example in the form of a flexible web or a belt, or inflexible, for example in the form of a drum. A drum can have a hollow cylindrical structure that provides for attachment of the drum to a drive that rotates the drum during the imaging process. Typically, a flexible electrically conductive substrate comprises an electrically insulating substrate and a thin layer of electrically conductive material onto which the photoconductive material is applied.

[0029] The electrically insulating substrate may be paper or a film forming polymer such as polyester (e.g., polyethylene terephthalate or polyethylene naphthalate), polyimide, polysulfone, polypropylene, nylon, polyester, polycarbonate, polyvinyl resin, polyvinyl fluoride, polystyrene and the like. Specific examples of polymers for supporting sub-

strates included, for example, polyethersulfone (Stabar™ S-100, available from ICI), polyvinyl fluoride (Tedlar®, available from E.I. DuPont de Nemours & Company), polybisphenol-A polycarbonate (Makrofol™, available from Mobay Chemical Company) and amorphous polyethylene terephthalate (Melinar™, available from ICI Americas, Inc.). The electrically conductive materials may be graphite, dispersed carbon black, iodide, conductive polymers such as polypyrroles and Calgon® conductive polymer 261 (commercially available from Calgon Corporation, Inc., Pittsburgh, Pa.), metals such as aluminum, titanium, chromium, brass, gold, copper, palladium, nickel, or stainless steel, or metal oxide such as tin oxide or indium oxide. In embodiments of particular interest, the electrically conductive material is aluminum. Generally, the photoconductor substrate has a thickness adequate to provide the required mechanical stability. For example, flexible web substrates generally have a thickness from about 0.01 to about 1 mm, while drum substrates generally have a thickness of from about 0.5 mm to about 2 mm.

[0030] The charge generating compound is a material which is capable of absorbing light to generate charge carriers, such as a dye or pigment. Non-limiting examples of suitable charge generating compounds include, for example, metal-free phthalocyanines (e.g., ELA 8034 metal-free phthalocyanine available from H.W. Sands, Inc. or Sanyo Color Works, Ltd., CGM-X01), metal phthalocyanines such as titanium phthalocyanine, copper phthalocyanine, oxytitanium phthalocyanine (also referred to as titanyl oxyphthalocyanine, and including any crystalline phase or mixtures of crystalline phases that can act as a charge generating compound), hydroxygallium phthalocyanine, squarylium dyes and pigments, hydroxy-substituted squarylium pigments, perylimides, polynuclear quinones available from Allied Chemical Corporation under the tradename Indofast® Double Scarlet, Indofast® Violet Lake B, Indofast® Brilliant Scarlet and Indofast® Orange, quinacridones available from DuPont under the tradename Monastral™ Red, Monastral™ Violet and Monastral™ Red Y, naphthalene 1,4,5,8-tetracarboxylic acid derived pigments including the perinones, tetrabenzoporphyrins and tetranaphthaloporphyrins, indigo- and thioindigo dyes, benzothioxanthene-derivatives, perylene 3,4,9,10-tetracarboxylic acid derived pigments, polyazo-pigments including bisazo-, trisazo- and tetrakisazo-pigments, polymethine dyes, dyes containing quinazoline groups, tertiary amines, amorphous selenium, selenium alloys such as selenium-tellurium, selenium-tellurium-arsenic and selenium-arsenic, cadmium sulphoselenide, cadmium selenide, cadmium sulphide, and mixtures thereof. For some embodiments, the charge generating compound comprises oxytitanium phthalocyanine (e.g., any phase thereof), hydroxygallium phthalocyanine or a combination thereof.

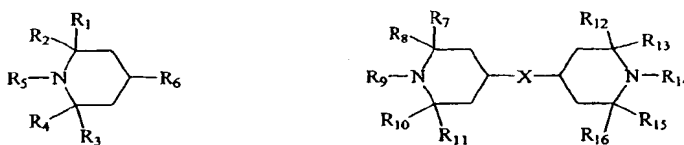
[0031] The photoconductive layer of this invention may contain an electron transport compound. Generally, any electron transport compound known in the art can be used. Non-limiting examples of suitable electron transport compound include, for example, bromoaniline, tetracyanoethylene, tetracyanoquinodimethane, 2,4,7-trinitro-9-fluorenone, 2,4,5,7-tetrinitro-9-fluorenone, 2,4,5,7-tetranitroxanthone, 2,4,8-trinitrothioxanthone, 2,6,8-trinitro-indeno[4H]-indeno[1,2-b]thiophene-4-one, and 1,3,7-trinitrodibenzo thiophene-5,5-dioxide, (2,3-diphenyl-1-indenylidene)malononitrile, 4H-thiopyran-1,1-dioxide and its derivatives such as 4-dicyanomethylene-2,6-diphenyl-4H-thiopyran-1,1-dioxide, 4-dicyanomethylene-2,6-di-m-tolyl-4H-thiopyran-1,1-dioxide, and unsymmetrically substituted 2,6-diaryl-4H-thiopyran-1,1-dioxide such as 4H-1,1-dioxo-2-(p-isopropylphenyl)-6-phenyl-4-(dicyanomethylidene)thiopyran and 4H-1,1-dioxo-2-(p-isopropylphenyl)-6-(2-thienyl)-4-(dicyanomethylidene)thiopyran, derivatives of phospho-2,5-cyclohexadiene, alkoxycarbonyl-9-fluorenylidene)malononitrile derivatives such as (4-n-butoxycarbonyl-9-fluorenylidene)malononitrile, (4-phenethoxycarbonyl-9-fluorenylidene)malononitrile, (4-carbitoxy-9-fluorenylidene)malononitrile, and diethyl(4-n-butoxy carbonyl-2,7-dinitro-9-fluorenylidene)-malonate, anthraquinodimethane derivatives such as 11,11,12,12-tetracyano-2-alkylanthraquinodimethane and 11,11-dicyano-12,12-bis(ethoxycarbonyl)anthraquinodimethane, anthrone derivatives such as 1-chloro-10-[bis(ethoxycarbonyl)methylene]anthrone, 1,8-dichloro-10-[bis(ethoxycarbonyl)methylene]anthrone, 1,8-dihydroxy-10-[bis(ethoxycarbonyl)methylene]anthrone, and 1-cyano-10-[bis(ethoxycarbonyl)methylene]anthrone, 7-nitro-2-aza-9-fluorenylidene-malononitrile, diphenoquinone derivatives, benzoquinone derivatives, naphthoquinone derivatives, quinine derivatives, tetracyanoethylenecyanoethylene, 2,4,8-trinitro thioxanthone, dinitrobenzene derivatives, dinitroanthracene derivatives, dinitroacridine derivatives, nitroanthraquinone derivatives, dinitroanthraquinone derivatives, succinic anhydride, maleic anhydride, dibromo maleic anhydride, pyrene derivatives, carbazole derivatives, hydrazone derivatives, N,N-dialkylaniline derivatives, diphenylamine derivatives, triphenylamine derivatives, triphenylmethane derivatives, tetracyano quinoedimethane, 2,4,5,7-tetrinitro-9-fluorenone, 2,4,7-trinitro-9-dicyanomethylene fluorenone, 2,4,5,7-tetranitroxanthone derivatives, and 2,4,8-trinitrothioxanthone derivatives. In some embodiments of interest, the electron transport compound comprises an (alkoxycarbonyl-9-fluorenylidene)malononitrile derivative, such as (4-n-butoxycarbonyl-9-fluorenylidene)malononitrile.

[0032] An electron transport compound and a UV light stabilizer can have a synergistic relationship for providing desired electron flow within the photoconductor. The presence of the UV light stabilizers alters the electron transport properties of the electron transport compounds to improve the electron transporting properties of the composite. UV light stabilizers can be ultraviolet light absorbers or ultraviolet light inhibitors that trap free radicals.

[0033] UV light absorbers can absorb ultraviolet radiation and dissipate it as heat. UV light inhibitors are thought to trap free radicals generated by the ultraviolet light and after trapping of the free radicals, subsequently to regenerate active stabilizer moieties with energy dissipation. In view of the synergistic relationship of the UV stabilizers with electron transport compounds, the particular advantages of the UV stabilizers may not be their UV stabilizing abilities, although

the UV stabilizing ability may be further advantageous in reducing degradation of the organophotoreceptor over time. While not wanting to be limited by theory, the synergistic relationship contributed by the UV stabilizers may be related to the electronic properties of the compounds, which contribute to the UV stabilizing function, by further contributing to the establishment of electron conduction pathways in combination with the electron transport compounds. In particular, the organophotoreceptors with a combination of the electron transport compound and the UV stabilizer can demonstrate a more stable acceptance voltage V_{acc} with cycling. The improved synergistic performance of organophotoreceptors with layers comprising both an electron transport compound and a UV stabilizer are described further in copending U.S. Patent Application Serial Number 10/425,333 filed on April 28, 2003 to Zhu, entitled "Organophotoreceptor With A Light Stabilizer," incorporated herein by reference.

[0034] Non-limiting examples of suitable light stabilizer include, for example, hindered trialkylamines such as Tinuvin 144 and Tinuvin 292 (from Ciba Specialty Chemicals, Terrytown, NY), hindered alkoxydialkylamines such as Tinuvin 123 (from Ciba Specialty Chemicals), benzotriazoles such as Tinuvan 328, Tinuvin 900 and Tinuvin 928 (from Ciba Specialty Chemicals), benzophenones such as Sanduvor 3041 (from Clariant Corp., Charlotte, N.C.), nickel compounds such as Arbostab (from Robinson Brothers Ltd, West Midlands, Great Britain), salicylates, cyanocinnamates, benzylidene malonates, benzoates, oxanilides such as Sanduvor VSU (from Clariant Corp., Charlotte, N.C.), triazines such as Cyagard UV-1164 (from Cytec Industries Inc., N.J.), polymeric sterically hindered amines such as Luchem (from Atochem North America, Buffalo, NY). In some embodiments, the light stabilizer is selected from the group consisting of hindered trialkylamines having the following formula:



where $R_1, R_2, R_3, R_4, R_6, R_7, R_8, R_{10}, R_{11}, R_{12}, R_{13}, R_{15}$ and R_{16} are, independently, hydrogen, alkyl group, or ester, or ether group; and R_5, R_9 , and R_{14} are, independently, alkyl group; and X is a linking group selected from the group consisting of $-O-CO-(CH_2)_m-CO-O-$ where m is between 2 to 20.

[0035] The binder generally is capable of dispersing or dissolving the charge transport compound (in the case of the charge transport layer or a single layer construction), the charge generating compound (in the case of the charge generating layer or a single layer construction) and/or an electron transport compound for appropriate embodiments. Examples of suitable binders for both the charge generating layer and charge transport layer generally include, for example, polystyrene-co-butadiene, polystyrene-co-acrylonitrile, modified acrylic polymers, polyvinyl acetate, styrene-alkyd resins, soya-alkyl resins, polyvinylchloride, polyvinylidene chloride, polyacrylonitrile, polycarbonates, polyacrylic acid, polyacrylates, polymethacrylates, styrene polymers, polyvinyl butyral, alkyd resins, polyamides, polyurethanes, polyesters, polysulfones, polyethers, polyketones; phenoxy resins, epoxy resins, silicone resins, polysiloxanes, poly(hydroxyether) resins, polyhydroxystyrene resins, novolak, poly(phenylglycidyl ether)-co-dicyclopentadiene, copolymers of monomers used in the above-mentioned polymers, and combinations thereof. Suitable binders include, for example, polyvinyl butyral, such as BX-1 and BX-5 from Sekisui Chemical Co. Ltd., Japan.

[0036] Suitable optional additives for any one or more of the layers include, for example, antioxidants, coupling agents, dispersing agents, curing agents, surfactants and combinations thereof.

[0037] The photoconductive element overall typically has a thickness from about 10 to about 45 microns. In the dual layer embodiments having a separate charge generating layer and a separate charge transport layer, charge generation layer generally has a thickness from about 0.5 to about 2 microns, and the charge transport layer has a thickness from about 5 to about 35 microns. In embodiments in which the charge transport compound and the charge generating compound are in the same layer, the layer with the charge generating compound and the charge transport composition generally has a thickness from about 7 to about 30 microns. In embodiments with a distinct electron transport layer, the electron transport layer has an average thickness from about 0.5 microns to about 10 microns and in further embodiments from about 1 micron to about 3 microns. In general, an electron transport overcoat layer can increase mechanical abrasion resistance, increases resistance to carrier liquid and atmospheric moisture, and decreases degradation of the photoreceptor by corona gases. A person of ordinary skill in the art will recognize that additional ranges of thickness within the explicit ranges above are contemplated and are within the present disclosure.

[0038] Generally, for the organophotoreceptors described herein, the charge generation compound is in an amount from about 0.5 to about 25 weight percent in further embodiments in an amount from about 1 to about 15 weight percent and in other embodiments in an amount from about 2 to about 10 weight percent, based on the weight of the photoconductive layer. The charge transport compound is in an amount from about 10 to about 80 weight percent, based

on the weight of the photoconductive layer, in further embodiments in an amount from about 35 to about 60 weight percent, and in other embodiments from about 45 to about 55 weight percent, based on the weight of the photoconductive layer. The optional electron transport compound, when present, can be in an amount of at least about 2 weight percent, in other embodiments from about 2.5 to about 25 weight percent, based on the weight of the photoconductive layer, and in further embodiments in an amount from about 4 to about 20 weight percent, based on the weight of the photoconductive layer. The binder is in an amount from about 15 to about 80 weight percent, based on the weight of the photoconductive layer, and in further embodiments in an amount from about 20 to about 75 weight percent, based on the weight of the photoconductive layer. A person of ordinary skill in the art will recognize that additional ranges within the explicit ranges of compositions are contemplated and are within the present disclosure.

[0039] For the dual layer embodiments with a separate charge generating layer and a charge transport layer, the charge generation layer generally comprises a binder in an amount from about 10 to about 90 weight percent, in further embodiments from about 15 to about 80 weight percent and in some embodiments in an amount of from about 20 to about 75 weight percent, based on the weight of the charge generation layer. The optional electron transport compound in the charge generating layer, if present, generally can be in an amount of at least about 2.5 weight percent, in further embodiments from about 4 to about 30 weight percent and in other embodiments in an amount from about 10 to about 25 weight percent, based on the weight of the charge generating layer. The charge transport layer generally comprises a binder in an amount from about 20 weight percent to about 70 weight percent and in further embodiments in an amount from about 30 weight percent to about 50 weight percent. A person of ordinary skill in the art will recognize that additional ranges of binder concentrations for the dual layer embodiments within the explicit ranges above are contemplated and are within the present disclosure.

[0040] For the embodiments with a single layer having a charge generating compound and a charge transport compound, the photoconductive layer generally comprises a binder, a charge transport compound and a charge generation compound. The charge generation compound can be in an amount from about 0.05 to about 25 weight percent and in further embodiment in an amount from about 2 to about 15 weight percent, based on the weight of the photoconductive layer. The charge transport compound can be in an amount from about 10 to about 80 weight percent, in other embodiments from about 25 to about 65 weight percent, in additional embodiments from about 30 to about 60 weight percent and in further embodiments in an amount of from about 35 to about 55 weight percent, based on the weight of the photoconductive layer, with the remainder of the photoconductive layer comprising the binder, and optionally additives, such as any conventional additives. A single layer with a charge transport composition and a charge generating compound generally comprises a binder in an amount from about 10 weight percent to about 75 weight percent, in other embodiments from about 20 weight percent to about 60 weight percent, and in further embodiments from about 25 weight percent to about 50 weight percent. Optionally, the layer with the charge generating compound and the charge transport compound may comprise an electron transport compound. The optional electron transport compound, if present, generally can be in an amount of at least about 2.5 weight percent, in further embodiments from about 4 to about 30 weight percent and in other embodiments in an amount from about 10 to about 25 weight percent, based on the weight of the photoconductive layer. A person of ordinary skill in the art will recognize that additional composition ranges within the explicit compositions ranges for the layers above are contemplated and are within the present disclosure.

[0041] In general, any layer with an electron transport layer can advantageously further include a UV light stabilizer. In particular, the electron transport layer generally can comprise an electron transport compound, a binder and an optional UV light stabilizer. An overcoat layer comprising an electron transport compound is described further in co-pending U.S. Patent Application Serial No. 10/396,536 to Zhu et al. entitled, "Organophotoreceptor With An Electron Transport Layer," incorporated herein by reference. For example, an electron transport compound as described above may be used in the release layer of the photoconductors described herein. The electron transport compound in an electron transport layer can be in an amount from about 10 to about 50 weight percent, and in other embodiments in an amount from about 20 to about 40 weight percent, based on the weight of the electron transport layer. A person of ordinary skill in the art will recognize that additional ranges of compositions within the explicit ranges are contemplated and are within the present disclosure.

[0042] The UV light stabilizer, if present, in any of one or more appropriate layers of the photoconductor generally is in an amount from about 0.5 to about 25 weight percent and in some embodiments in an amount from about 1 to about 10 weight percent, based on the weight of the particular layer. A person of ordinary skill in the art will recognize that additional ranges of compositions within the explicit ranges are contemplated and are within the present disclosure.

[0043] For example, the photoconductive layer may be formed by dispersing or dissolving the components, such as one or more of a charge generating compound, a charge transport compound, an electron transport compound, a UV light stabilizer, and a polymeric binder in organic solvent, coating the dispersion and/or solution on the respective underlying layer and drying the coating. In particular, the components can be dispersed by high shear homogenization, ball-milling, attritor milling, high energy bead (sand) milling or other size reduction processes or mixing means known in the art for effecting particle size reduction in forming a dispersion.

[0044] The photoreceptor may optionally have one or more additional layers as well. An additional layer can be, for example, a sub-layer or an overcoat layer, such as a barrier layer, a release layer, a protective layer, or an adhesive layer. A release layer or a protective layer may form the uppermost layer of the photoconductor element. A barrier layer may be sandwiched between the release layer and the photoconductive element or used to overcoat the photoconductive element. The barrier layer provides protection from abrasion to the underlayers. An adhesive layer locates and improves the adhesion between a photoconductive element, a barrier layer and a release layer, or any combination thereof. A sub-layer is a charge blocking layer and locates between the electrically conductive substrate and the photoconductive element. The sub-layer may also improve the adhesion between the electrically conductive substrate and the photoconductive element.

[0045] Suitable barrier layers include, for example, coatings such as crosslinkable siloxanol-colloidal silica coating and hydroxylated silsesquioxane-colloidal silica coating, and organic binders such as polyvinyl alcohol, methyl vinyl ether/maleic anhydride copolymer, casein, polyvinyl pyrrolidone, polyacrylic acid, gelatin, starch, polyurethanes, polyimides, polyesters, polyamides, polyvinyl acetate, polyvinyl chloride, polyvinylidene chloride, polycarbonates, polyvinyl butyral, polyvinyl acetoacetal, polyvinyl formal, polyacrylonitrile, polymethyl methacrylate, polyacrylates, polyvinyl carbazoles, copolymers of monomers used in the above-mentioned polymers, vinyl chloride/vinyl acetate/vinyl alcohol terpolymers, vinyl chloride/vinyl acetate/maleic acid terpolymers, ethylene/vinyl acetate copolymers, vinyl chloride/vinylidene chloride copolymers, cellulose polymers, and mixtures thereof. The above barrier layer polymers optionally may contain small inorganic particles such as fumed silica, silica, titania, alumina, zirconia, or a combination thereof. Barrier layers are described further in U.S. Patent 6,001,522 to Woo et al., entitled "Barrier Layer For Photoconductor Elements Comprising An Organic Polymer And Silica," incorporated herein by reference. The release layer topcoat may comprise any release layer composition known in the art. In some embodiments, the release layer is a fluorinated polymer, siloxane polymer, fluorosilicone polymer, silane, polyethylene, polypropylene, polyacrylate, or a combination thereof. The release layers can comprise crosslinked polymers.

[0046] The release layer may comprise, for example, any release layer composition known in the art. In some embodiments, the release layer comprises a fluorinated polymer, siloxane polymer, fluorosilicone polymer, polysilane, polyethylene, polypropylene, polyacrylate, poly(methyl methacrylate-co-methacrylic acid), urethane resins, urethane-epoxy resins, acrylated-urethane resins, urethane-acrylic resins, or a combination thereof. In further embodiments, the release layers comprise crosslinked polymers.

[0047] The protective layer can protect the organophotoreceptor from chemical and mechanical degradation. The protective layer may comprise any protective layer composition known in the art. In some embodiments, the protective layer is a fluorinated polymer, siloxane polymer, fluorosilicone polymer, polysilane, polyethylene, polypropylene, polyacrylate, poly(methyl methacrylate-co-methacrylic acid), urethane resins, urethane-epoxy resins, acrylated-urethane resins, urethane-acrylic resins, or a combination thereof. In some embodiments of particular interest, the release layers are crosslinked polymers.

[0048] An overcoat layer may comprise an electron transport compound as described further in copending U.S. Patent Application Serial No. 10/396,536, filed on March 25, 2003 to Zhu et al. entitled, "Organoreceptor With An Electron Transport Layer," incorporated herein by reference. For example, an electron transport compound, as described above, may be used in the release layer of this invention. The electron transport compound in the overcoat layer can be in an amount from about 2 to about 50 weight percent, and in other embodiments in an amount from about 10 to about 40 weight percent, based on the weight of the release layer. A person of ordinary skill in the art will recognize that additional ranges of composition within the explicit ranges are contemplated and are within the present disclosure.

[0049] Generally, adhesive layers comprise a film forming polymer, such as polyester, polyvinylbutyral, polyvinylpyrrolidone, polyurethane, polymethyl methacrylate, poly(hydroxy amino ether) and the like.

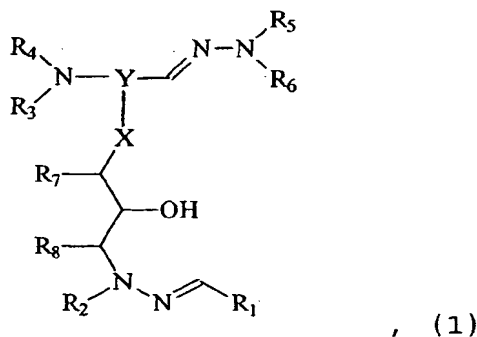
[0050] Sub-layers can comprise, for example, polyvinylbutyral, organosilanes, hydrolyzable silanes, epoxy resins, polyesters, polyamides, polyurethanes, silicones and the like. In some embodiments, the sub-layer has a dry thickness between about 20 Angstroms and about 2,000 Angstroms. Sublayers containing metal oxide conductive particles can be between about 1 and about 25 microns thick. A person of ordinary skill in the art will recognize that additional ranges of compositions and thickness within the explicit ranges are contemplated and are within the present disclosure.

[0051] The charge transport compounds as described herein, and photoreceptors including these compounds, are suitable for use in an imaging process with either dry or liquid toner development. For example, any dry toners and liquid toners known in the art may be used in the process and the apparatus of this invention. Liquid toner development can be desirable because it offers the advantages of providing higher resolution images and requiring lower energy for image fixing compared to dry toners. Examples of suitable liquid toners are known in the art. Liquid toners generally comprise toner particles dispersed in a carrier liquid. The toner particles can comprise a colorant/pigment, a resin binder, and/or a charge director. In some embodiments of liquid toner, a resin to pigment ratio can be from 1:1 to 10:1, and in other embodiments, from 4:1 to 8:1. Liquid toners are described further in Published U.S. Patent Applications 2002/0128349, entitled "Liquid Inks Comprising A Stable Organosol," 2002/0086916, entitled "Liquid Inks Comprising Treated Colorant Particles," and 2002/0197552, entitled "Phase Change Developer For Liquid Electrophotography,"

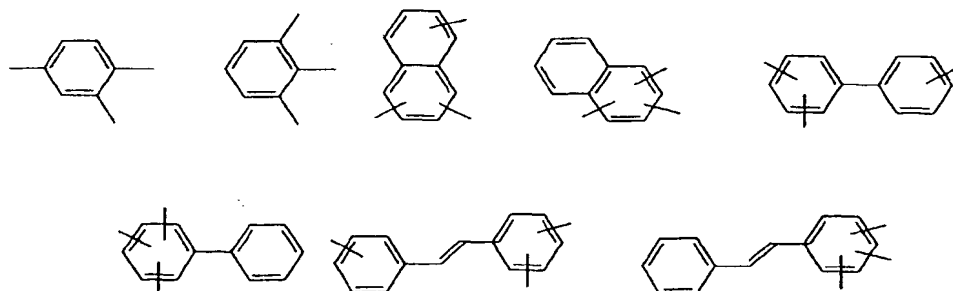
all three of which are incorporated herein by reference.

Charge Transport Compound

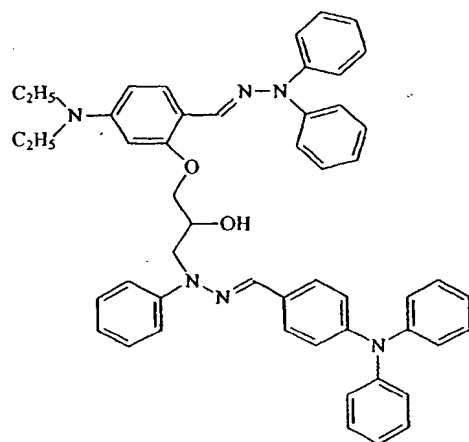
[0052] As described herein, an organophotoreceptor comprises a charge transport compound having the formula



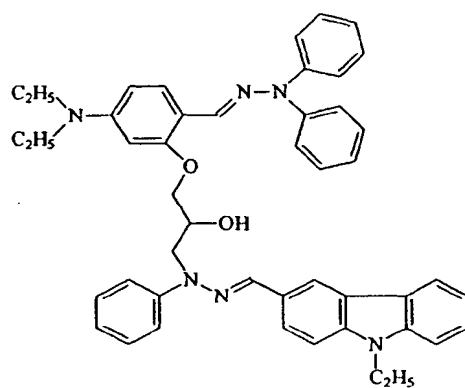
where R_1 is a carbazole group, a julolidine group, or a p-(N,N-disubstituted)arylamine, R_2 , R_3 , R_4 , R_5 and R_6 are, independently, hydrogen, an alkyl group or an aryl group, R_7 and R_8 are, independently, hydrogen, an alkyl group, or an aryl group, X is oxygen, sulfur, or a NR' group where R' is hydrogen, an alkyl, or an aryl group, and Y is an aryl group. As noted in formula (1), Y is at least trivalent. Non-limiting examples of trivalent aryl groups that are suitable as Y have the following formulae:



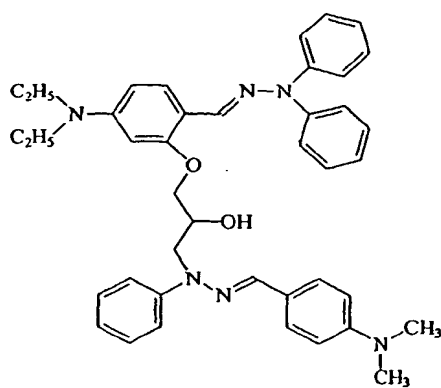
[0053] Specific, non-limiting examples of suitable charge transport compounds within the general Formula (1) of the present invention have the following structures.



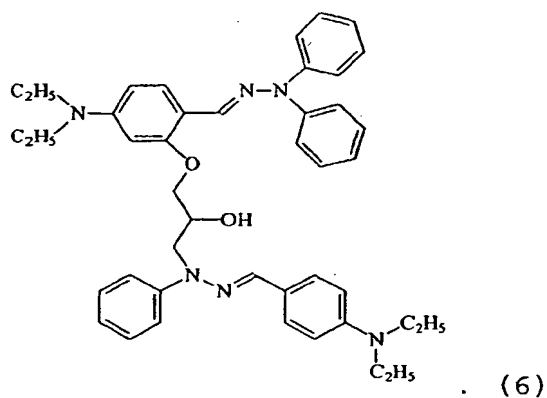
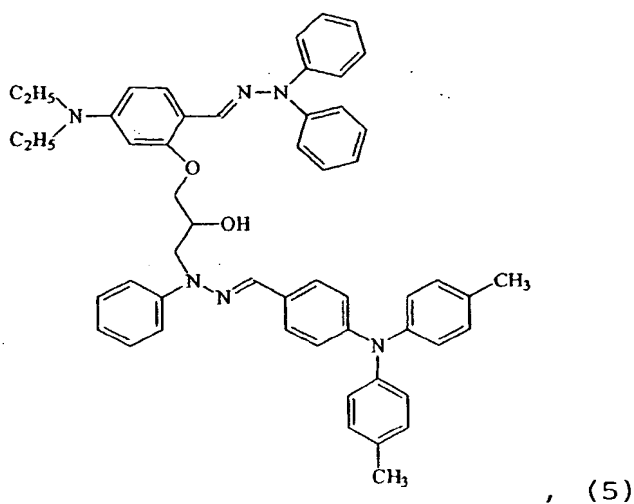
, (2)



, (3)

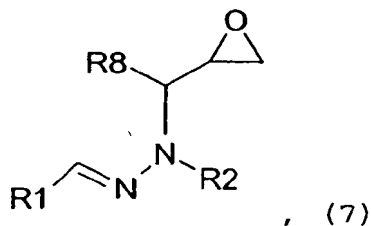


, (4)

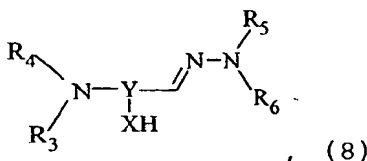


Synthesis of Charge Transport Compounds

[0054] In general, the charge transport compounds can be prepared as two precursors, each having a hydrazone group that are then reacted to form the charge transport compound. Specifically, one precursor can be formed with the structure



where R₁, R₂ and R₈ are selected to yield the desired specific structure within the general structure of formula (1) above. The second precursor has the structure



where R₃, R₄, R₅, R₆, and X are selected to yield the desired specific structure within the general framework of formula (1) above. The active hydrogen on the X group adds to the epoxy group to form the product under alkaline catalysis for an epoxy reaction.

Synthesis of First Precursor Of Formula (7)

[0055] The charge transport compounds with a hydrazone bonded to the epoxy group generally are synthesized by forming the desired substituted hydrazone, which is reacted at the secondary amine to form the epoxy group. In particular, the aromatic-substituted secondary amine reacts with the epichlorhydrin by way of the active hydrogen of the secondary amine in a base catalyzed reaction to form the epoxy group with a -CH₂- group between the epoxy group and the amine. The hydrazone is formed from the reaction of an aryl substituted hydrazine with an aldehyde or ketone having a N,N-disubstituted arylamine, as describe further below. Suitable hydrazines and arylamines are described further below.

[0056] The aromatic-substituted hydrazine supplies the R₂ group from formula (7) above, and an N,N-disubstituted amino aryl substituted aldehyde or ketone supplies the R₁ group of formula (7). In the reaction of the aldehyde or ketone with the hydrazine, the oxygen of the aldehyde/ketone group is replaced with the double bonded carbon.

Hydrazines

[0057] The synthesis of some representative hydrazines is described in the following.

1,1-Dinaphthylhydrazine

[0058] 1,1-Dinaphthylhydrazine can be prepared according to the procedure described in Staschkow, L.I.; Matevosyan, R.O. Journal of the General Chemistry (1964) 34, 136, which is incorporated herein by reference. A suspension of 0.07 mole of the naphthyl nitrosamine in 750 ml of ether is cooled to 5-8 °C and treated with 150 g of zinc dust. Acetic acid (70 ml) is then added drop wise with stirring. To complete the reaction, 40g of zinc dust is added. The reaction mixture is heated and filtered from the sludge. The mother liquor is washed with 10% sodium carbonate solution and dried with solid KOH. The ether is distilled off to give the crystalline hydrazine, which is crystallized from ethanol or butanol. Other symmetric disubstituted hydrazines can be synthesized based on an equivalent process.

N-Phenyl-N-sulfolan-3-ylhydrazine

[0059] N-Phenyl-N-sulfolan-3-ylhydrazine can be prepared according to the procedure described in Great Britain Patent No. 1,047,525 by Mason, which is incorporated herein by reference. To a mixture of 0.5 mole of butadiene sulfone (commercially available from Aldrich, Milwaukee, WI) and 0.55 mole of phenylhydrazine (commercially available from Aldrich, Milwaukee, WI) was added 0.005 mole 40% aqueous potassium hydroxide solution. The mixture was kept for 2 hours at 60°C whereupon a solid separated. After 10 hours the solid was filtered off to give N-phenyl-N-sulfolan-3-ylhydrazine (53%) having a melting point of 120-121°C (MeOH).

N-Pyrrol-2-yl-N-phenylhydrazine

[0060] N-Pyrrol-2-yl-N-phenylhydrazine can be prepared according to the procedure described in Japanese Patent No. 05148210 by Myamoto, incorporated herein by reference.

1-Phenyl-1-(1-benzyl-1H-tetrazol-5-yl)hydrazine

[0061] 1-Phenyl-1-(1-benzyl-1H-tetrazol-5-yl)hydrazine can be prepared according to the procedure described in Tetrahedron (1983), 39(15), 2599-608 by Atherton et al., incorporated herein by reference.

N-(4-Stilbenyl)-N-phenylhydrazine

[0062] N-(4-Stilbenyl)-N-phenylhydrazine can be prepared according to the procedure described in Zh. Org. Khim. (1967), 3(9), 1605-3 by Matevosyan et al., incorporated herein by reference. Following this procedure, to a mixture of phenylhydrazine (97 g, 0.9 mole, commercially available from Aldrich, Milwaukee, WI) and p-chlorostilbene (21.4 g, 0.1 mole, commercially available from Spectrum Quality Products, Inc., Gardena, CA; Web: www.spectrumchemical.com) heated to boiling temperature, sodium was slowly added until there was no more discharge of red coloration. After boiling for some time the mixture was dissolved in 1750 ml of ethanol and cooled to -15 °C. The precipitated product was recrystallized to give 28% of N-(4-stilbenyl)-N-phenylhydrazine.

N-(5-Benzotriazolyl)-N-phenylhydrazine

[0063] N-(5-benzotriazolyl)-N-phenylhydrazine can be prepared according to the procedure that follows. To a mixture of phenylhydrazine (97 g, 0.9 mole, commercially available from Aldrich, Milwaukee, WI) and 5-chlorobenzotriazole (15.4 g, 0.1 mole, commercially available from Aldrich, Milwaukee, WI) heated to boiling temperature, sodium is slowly added until there is no more discharge of red coloration. After boiling for some time the mixture is cooled to room temperature. The product is isolated and purified.

N-Phenyl-N-sulfolan-3-ylhydrazine

[0064] N-Phenyl-N-sulfolan-3-ylhydrazine can be prepared according to the procedure described in Great Britain Patent No. 1,047,525 by Mason, incorporated herein by reference. Following this procedure, to a mixture of 0.5 mole of butadiene sulfone (commercially available from Aldrich, Milwaukee, WI) and 0.55 mole of phenylhydrazine (commercially available from Aldrich, Milwaukee, WI), a 0.005 mole 40% aqueous potassium hydroxide solution was added. The mixture was kept for 2 hours at 60 °C whereupon a solid separated. After 10 hours the solid was filtered off to give N-phenyl-N-sulfolan-3-ylhydrazine (I) (93%) having a melting point of 119-20 °C (MeOH).

N-4-[(9H-fluoren-9-ylidene)benzyl]-N-phenylhydrazine

[0065] N-4-[(9H-fluoren-9-ylidene)benzyl]-N-phenylhydrazine can be prepared according to the procedure similar to that described in Zh. Org. Khim. (1967), 3(9), 1605-3 by Matevosyan et al., incorporated herein by reference. Following this procedure, to a mixture of phenylhydrazine (97 g, 0.9 mole, commercially available from Aldrich, Milwaukee, WI) and p-9-(4-chlorobenzylidene)fluorene (28.9 g, 0.1 mole, commercially available from Aldrich, Milwaukee, WI) heated to boiling temperature, sodium was slowly added until there was no more discharge of red coloration. After boiling for some time the mixture was dissolved in 1750 ml of ethanol and cooled to -15 °C. The precipitated product was recrystallized to give N-4-[(9H-fluoren-9-ylidene)benzyl]-N-phenylhydrazine.

5-Methyl-1-Phenyl-3-(1-Phenylhydrazino)-Pyrazole

[0066] 5-Methyl-1-phenyl-3-(1-phenylhydrazino)-pyrazole can be prepared according to the procedure described in J. Chem. Soc. C (1971), (12), 2314-17 by Boyd et al., incorporated herein by reference.

4-Methylsulfonylphenylhydrazine (Registry Number 877-66-7)

[0067] 4-Methylsulfonylphenylhydrazine is commercially available from Fisher Scientific USA, Pittsburgh, PA (1-800-766-7000).

1,1'-(Sulfonyldi-4,1-phenylene)bishydrazine (Registry Number 14052-65-4)

[0068] 1,1'-(Sulfonyldi-4,1-phenylene)bishydrazine dihydrochloride is commercially available from Vitas-M, Moscow, Russia; (Phone: 7 095 939 5737)

Arylaldehydes

[0069] Representative arylaldehydes for reacting with the hydrozones can be obtained as follows.

Synthesis Of Julolidine Aldehyde

[0070] Julolidine (100 g, 0.6 moles, commercially obtained from Aldrich Chemicals Co, Milwaukee, WI 53201) was dissolved in DMF (200 ml, commercially obtained from Aldrich) in a 500 ml three neck round bottom flask. The flask was cooled to 0°C in ice bath. POCl₃ (107 g, 0.7 mole, Aldrich) was added drop wise while keeping the temperature below 5 °C. After the addition of POCl₃ was completed, the flask was warmed to room temperature and placed in a steam bath while stirring for a period of 1 hour. The flask was cooled to room temperature and the solution was added slowly to a large excess of distilled water with good agitation. Stirring was continued for additional 2 hours. The solid was filtered off and washed repeatedly with water until the effluent water became neutral. The product was dried in vacuum oven at 50 C for 4 hours.

Other Aryl Aldehydes

[0071] Suitable commercially available (N,N-disubstituted)arylamine aldehydes are available from Aldrich (Milwaukee, WI) including, for example, diphenylamino-benzaldehyde ((C₆H₅)₂NC₆H₄CHO) and 9-ethyl-3-carbazolecarboxyaldehyde. Also, the synthesis of N-ethyl-3,6-diformylcarbazole is described below in the examples.

Synthesis of Hydrazones

[0072] A hydrazine can be reacted with an appropriate aromatic aldehyde/ketone to form a desired hydrazone charge transfer compound. The reactions can be catalyzed by an appropriate amount of concentrated acid, in particular sulfuric acid. After mixing in the catalytic amount of acid with the hydrazine and aromatic aldehyde, the mixture can be refluxed for about 2 hours to about 16 hours. The initial product can be purified by recrystallization. The synthesis of selected compounds from the formulas above are described below in the Examples, and the other compounds described herein can be similarly synthesized.

[0073] In some embodiments, the hydrazines may be obtained in an acidified hydrochloride form, as noted above. For these embodiments, the hydrazine hydrochloride can be reacted with an aqueous carbonate base while stirring the mixture. An excess of carbonate base can be added, such as 1.2 moles of potassium carbonate for embodiments with one mole of hydrazine hydrochloride per mole hydrazine or 2.4 moles of potassium carbonate for embodiments with one mole of hydrazine dihydrochloride per mole hydrazine. Some specific examples are presented below.

Synthesis of Second Precursor of Formula (8)

[0074] The synthesis of the second precursor can be performed by reacting a hydrazine with an aminoaryl aldehyde. The synthesis of suitable hydrazines is described above. Suitable aminoaryl aldehydes have a hydroxyl group, a thiol group or a second amino group to react with the first precursor. Some suitable aminoaryl aldehydes are commercially available. For example, 4-salicylaldehyde and 9-formyl-8-hydroxyjulolidine are available from Aldrich. The aminoaryl aldehydes can be reacted with the hydrazines, for example, as described for one embodiment in Example 1 below.

[0075] The invention will now be described further by way of the following illustrative and non-limiting examples. These examples are to be viewed as being illustrative of specific materials falling within the broader disclosure presented above and are not to be viewed as limiting the broader disclosure in any way.

EXAMPLESExample 1 - Synthesis of Charge Transfer Compound Precursors

[0076] This example described the synthesis of several intermediates used in the synthesis of several specific embodiments of the charge transfer compounds. The synthesis of the charge transfer compounds is described in the next Example.

Preparation of 4-(diethylamino)salicylaldehyde N,N-diphenylhydrazone.

[0077] This example described the formation of a reactant used in the formation of several charge transport compounds in the next example.

[0078] A solution of N,N-diphenylhydrazine hydrochloride (79.5 g, 0.36 mol, commercially available from Aldrich, Milwaukee, WI) in ethanol (500 ml) was slowly added to a solution of 4-(diethylamino)salicylaldehyde (58.0 g, 0.3 mol, commercially available from Aldrich, Milwaukee, WI) in ethanol (500 ml) to form a reaction mixture. The reaction mixture was refluxed until all aldehyde reacted, which took about 0.5 h. Then, the solvent was evaporated. After the evaporation

of the solvent (800 ml), the residue was treated with ether (ethyl ether). The ether extract was washed with water until the pH of the wash water was neutral. The separated organic liquid was dried over anhydrous magnesium sulphate, treated with activated charcoal and filtered. The solvent was then removed, and ethanol was added to the residue to crystallize the product. 4-(Diethylamino)salicylaldehyde N,N-diphenyl-hydrazone (85 g; 78.8 %) was filtered off and washed with ethanol. The obtained hydrazone was recrystallized from the mixture of 2-propanol and ether (10:1). The product was found to have a melting point of 95.5-96.5°C following recrystallization. A proton NMR spectrum was obtained for the product. The ¹H-NMR spectrum (100 MHz, CDCl₃) had the following peaks, ppm: 11.55 (s, 1H); 7.55-6.95 (m, 11H); 6.7 (d, J=8.6 Hz, 1H); 6.23 (s, 1H); 6.1 (d, J=8.6 Hz, 1H); 3.3 (q, J=8.0 Hz, 4H); 1.1 (t, J=8.0 Hz, 6H); where s is singlet, d is doublet, m is multiplet, q is quartet, t is triplet. An elemental analysis yielded, %: C 76.68; H 7.75; N 11.45. C₂₃H₂₅N₃O has a calculated elemental distribution of, %: C 76.85; H 7.01; N 11.69.

Preparation of 4-(Diphenylamino)benzaldehyde-N-2, 3-epoxypropyl-N-Phenylhydrazone

[0079] Phenylhydrazine (0.1 mole, commercially available from Aldrich, Milwaukee, WI) and 4-(Diphenylamino) benzaldehyde (0.1 mole, available from Fluka, Buchs SG, Switzerland) were dissolved in 100 ml of isopropanol in a 250 ml 3-neck round bottom flask equipped with reflux condenser and mechanical stirrer. The solution was refluxed for 2 hours. Thin layer chromatography indicated the disappearance of the starting materials. At the end of the reaction, the mixture was cooled to room temperature. The 4-(diphenylamino) benzaldehyde phenylhydrazone crystals that formed upon standing were filtered off and washed with isopropanol and dried in vacuum oven at 50°C for 6 hours.

[0080] A mixture of 4-(diphenylamino) benzaldehyde phenylhydrazone (3.6 g, 0.01 mole), 85% powdered potassium hydroxide (2.0 g, 0.03 mole) and anhydrous potassium carbonate in 25 ml of epichlorohydrin was stirred vigorously at 55-60°C for 1.5-2 hours. The course of the reaction was monitored using thin layer chromatography on silica gel 60 F254 plates (commercially available from Merck, Whitehouse Station, NJ) using 1:4 v/v mixture of acetone and hexane as eluant. After termination of the reaction, the mixture was cooled to room temperature, diluted with ether, and washed with water until the wash water had a neutral pH. The organic layer was dried over anhydrous magnesium sulfate, treated with activated charcoal and filtered. Ether was removed and the residue was dissolved in a 1:1 volume per volume mixture of toluene and isopropanol. The crystals formed upon standing were filtered off and washed with isopropanol to give 3.0 g of product (71.4% yield) with a melting point of 141-142.5°C. The product was recrystallized from a 1:1 mixture of toluene and isopropanol. The product was characterized with ¹H-NMR in CDCl₃ (250MHz instrument) with peaks observed at the following delta values in ppm: 7.65-6.98 (m, 19H), 6.93 (t, J= 7.2Hz, 1H), 4.35 (dd, 1H), 3.99 (dd, 1H), 3.26 (m, 1H), 2.84 (dd, 1H), 2.62 (dd, 1H). An elemental analysis yielded the following results in weight percent: %C = 80.02, %H = 6.31, %N = 9.91, which compares with calculated values for C₂₈H₂₅N₃O of %C = 80.16, %H = 6.01, %N = 10.02.

Preparation of 9-ethyl-3-carbazolecarboxaldehyde-N-2,3-epoxypropyl-N-phenyl hydrazone

[0081] Phenylhydrazine (0.1 mole, commercially available from Aldrich, Milwaukee, WI) and 9-ethyl-3-carbazolecarboxaldehyde (0.1 mole, available from Aldrich Chemical, Milwaukee, WI) were dissolved in 100 ml of isopropanol in 250 ml 3-neck round bottom flask equipped with a reflux condenser and a mechanical stirrer. The solution was refluxed for 2 hours. Thin layer chromatography indicated the disappearance of the starting materials. At the end of the reaction, the mixture was cooled to room temperature. The 9-ethyl-3-carbazolecarboxaldehyde phenylhydrazone crystals formed upon standing were filtered off and washed with isopropanol and dried in vacuum oven at 50°C for 6 hours.

[0082] A mixture of 9-ethyl-3-carbazolecarboxaldehyde phenylhydrazone (3.1 g, 0.01 mole), 85% powdered potassium hydroxide (2.0 g, 0.03 mole) and anhydrous potassium carbonate in 25 ml of epichlorohydrin was stirred vigorously at 55-60°C for 1.5-2 hours. The course of the reaction was monitored using thin layer chromatography on silica gel 60 F254 plates (commercially available from Merck) using 1:4 v/v mixture of acetone and hexane as eluant. After termination of the reaction, the mixture was cooled to room temperature, diluted with ether and washed with water until the wash water had a neutral pH. The organic layer was dried over anhydrous magnesium sulfate, treated with activated charcoal and filtered. Ether was removed and the residue was dissolved in a 1:1 mixture of toluene and isopropanol. The crystals formed upon standing were filtered off and washed with isopropanol to give 3.0 g of product (81.2% yield) with a melting point of 136-137 °C. The product was recrystallized from 1:1 mixture of toluene and isopropanol. The product was characterized with ¹H-NMR in CDCl₃ (250MHz) which yielded peaks at the following delta values in ppm: 8.35 (s, 1H), 8.14 (d, J=7.8Hz, 1H), 7.93 (d, J= 7.6Hz, 1H), 7.90 (s, 1H), 7.54-7.20 (m, 8H), 6.96 (t, J=7.2Hz, 1H), 4.37 (m, 3H), 4.04 (dd, J1=4.3 Hz, J2 = 16.4 Hz, 1H), 3.32 (m, 1H), 2.88 (dd, 1H), 2.69 (dd, 1H), 1.44 (t, J=7.2 Hz, 3H). Elemental analysis yielded the following results in weight percent %C=78.32, %H = 6.41, %N = 11.55; which compares with calculated values for C₂₄H₂₃N₃O of %C = 78.02, %H = 6.28, %N = 11.37.

Preparation of 4-dimethylaminobenzaldehyde-N-2, 3-epoxypropyl-N-phenylhydrazone

[0083] The preparation of 4-dimethylaminobenzaldehyde-N-2, 3-epoxypropyl-N-phenylhydrazone was performed as described above for 4-(diphenylamino)benzaldehyde-N-2, 3-epoxypropyl-N-phenylhydrazone except that 4-(dimethylamino)benzaldehyde (Aldrich) was used instead of 4-(Diphenylamino) benzaldehyde.

Preparation of 4-(4,4'-di(methylphenyl)amino) benzaldehyde - N-2,3-epoxypropyl-N-phenylhydrazone

[0084] The preparation of 4-(4,4'-di(methylphenyl)amino)benzaldehyde-N-2, 3-epoxypropyl-N-phenylhydrazone was performed as described above for 4-(diphenylamino)benzaldehyde-N-2, 3-epoxypropyl-N-phenylhydrazone except that 4-(4,4'-di(methylphenyl)amino) benzaldehyde (Aldrich) was used instead of 4-(Diphenylamino) benzaldehyde.

Preparation of 4-diethylaminobenzaldehyde-N-2, 3-epoxypropyl-N-Phenylhydrazone

[0085] The preparation of 4-diethylaminobenzaldehyde-N-2, 3-epoxypropyl-N-phenylhydrazone was performed as described above for 4-(diphenylamino)benzaldehyde-N-2, 3-epoxypropyl-N-phenylhydrazone except that 4-(diethylamino)benzaldehyde (Aldrich) was used instead of 4-(Diphenylamino) benzaldehyde.

Example 2 - Synthesis of Charge Transport Compounds

[0086] This example describes the synthesis of five charge transport compounds corresponding to formulas 2-6 above.

Compound 2

[0087] A 0.6 ml (4.45 mmol) quantity of triethylamine (TEA) was added to a mixture of 4-(diethylamino)salicylaldehyde N,N-diphenylhydrazone (4.0 g, 11.13 mmol) and 4-(diphenylamino)benzaldehyde N-2,3-epoxypropyl-N-phenylhydrazone (4.67 g, 11.13 mmol) and 15 ml of 2-butanone. The mixture was refluxed until one of the starting compounds disappeared, which required 38 hours, as determined by thin layer chromatography (TLC) using silica gel 60 F254 plates, commercially available from Merck, and 1:4 v/v mixture of acetone and hexane as eluent. At the end of the reaction, 2-butanone and TEA were distilled off and the residue was subjected to chromatography using a column packed with silica gel (Merck grade 9385, commercially obtained from Aldrich, Milwaukee, WI) and a 4:1 v/v solution of hexane and acetone as the eluant. The product was crystallized from a mixture of 2-propanol and ether (10:1). The crystals formed were filtered off and washed with the mixture of 2-propanol and *n*-hexane (1:1) to give 6.5 g (74.7 %) of Compound 2. The product had a melting point of 161-162 °C after crystallization from 10:1 v/v of 2-propanol and ether. A ¹HNMR spectrum (250 MHz, CDCl₃) yielded the following peaks, ppm: 7.85 (d, *J*=8.8 Hz; 1H); 7.60-6.92 (m, 31H); 6.37 (d, *J*=8.8 Hz; 1H); 6.02 (s, 1H); 4.18 (m, 1H); 4.03-3.68 (m, 4H); 3.31 (q, *J*=7.1 Hz; 4H); 2.64 (d, *J*=6.5 Hz; 1H); 1.11 (t, *J*=7.1 Hz; 6H). An elemental analysis yielded, %: C 78.44; H 6.29; N 10.61. C₅₁H₅₀N₆O₂ has a calculated elemental distribution of %: C 78.64; H 6.47; N 10.79.

Compound 3

[0088] Compound 3 was prepared and isolated following the procedure used to prepare Compound 2 except that instead of 4-(diphenylamino)benzaldehyde N-2,3-epoxypropyl-N-phenylhydrazone, 9-ethyl-3-carbazolecarboxaldehyde N-2,3-epoxypropyl-N-phenylhydrazone (4.11 g, 11.13 mmol) was used. The yield was 5.9 g (72.8 %) and the product had a melting point of 108.5-109.5 °C after crystallization from 10:1 v/v of 2-propanol and ether. A ¹HNMR spectrum (250 MHz, CDCl₃) had peaks at, ppm: 8.17 (s, 1H); 8.12 (d, *J*=7.5 Hz; 1H); 7.94 (d, *J*=8.8 Hz; 1H); 7.85 (d, *J*=7.6 Hz; 1H); 7.74 (s, 1H); 7.61 (s, 1H); 7.55-6.96 (m, 19H); 6.38 (d, *J*=8.8 Hz; 1H); 6.02 (s, 1H); 4.37 (q, *J*=7.3 Hz; 2H); 4.25 (m, 1H); 4.03-3.72 (m, 4H); 3.26 (q, *J*=7.1 Hz; 4H); 2.72 (d, *J*=7.2 Hz; 1H); 1.44 (t, *J*=7.1 Hz; 3H); 1.06 (t, *J*=7.1 Hz; 6H). An elemental analysis yielded, %: C 77.21; H 6.48; N 11.68. C₄₇H₄₈N₆O₂ has a calculated elemental distribution of, %: C 77.44; H 6.64; N 11.53.

Compound 4

[0089] Compound 4 was prepared following the procedure used to prepare Compound 2 except that instead of 4-(diphenylamino)benzaldehyde N-2,3-epoxypropyl-N-phenylhydrazone, 4-dimethylaminobenzaldehyde N-2,3-epoxypropyl-N-phenylhydrazone (3.29 g, 11.13 mmol) was used. The reaction time was 34 h. After removal of the solvent, the residue was crystallized from a mixture of toluene and 2-propanol. The yield was 4.8 g (65.8 %), and the product

had a melting point of 159.5-160.5 °C when crystallized from the mixture of toluene and 2-propanol. A ¹HNMR spectrum (250 MHz, CDCl₃) yielded peaks at, ppm: 7.88 (d, *J*=8.8 Hz; 1H); 7.55 (s, 1H); 7.48 (m, 3H); 7.36-6.92 (m, 17H); 6.69 (d, *J*=8.9 Hz; 2H); 6.38 (d, *J*=8.8 Hz; 1H); 6.02 (s, 1H); 4.17 (m, 1H); 3.99-3.62 (m, 4H); 3.31 (q, *J*=7.1 Hz; 4H); 2.98 (s, 6H); 2.79 (d, *J*=6.5 Hz; 1H); 1.12 (t, *J*=7.1 Hz; 6H). An elemental analysis yielded, %: C 75.01; H 6.91; N 12.68. C₄₁H₄₆N₆O₂ has a calculated elemental distribution of, %: C 75.20; H 7.08; N 12.83.

Compound 5

[0090] Compound 5 was prepared and isolated following the procedure used to prepare Compound 2 except that instead of 4-(diphenylamino)benzaldehyde N-2,3-epoxypropyl-N-phenylhydrazone, 4-(4,4'-dimethyldiphenylamino)benzaldehyde N-2,3-epoxypropyl-N-phenylhydrazone (4.98 g, 11.13 mmol) was used. Compound 5 was obtained as an oil at a yield of 6.8 g (76.4 %). A ¹HNMR spectrum (100 MHz, CDCl₃) yielded peaks at, ppm: 7.9 (d, *J*=8.8 Hz; 1H); 7.7-6.9 (m, 29H); 6.4 (d, *J*=8.8 Hz; 1H); 6.05 (s, 1H); 4.3 (m, 1H); 4.1-3.7 (m, 4H); 3.35 (q, *J*=7.1 Hz; 4H); 2.65 (m, 1H); 2.4 (s, 6H); 1.15 (t, *J*=7.1 Hz; 6H). AN elemental analysis yielded, %: C 78.69; H 6.58; N 10.60. C₅₃H₅₄N₆O₂ has a calculated elemental distribution of, %: C 78.88; H 6.74; N 10.41.

Compound 6

[0091] Compound 6 was prepared and isolated following the procedure used to prepare Compound 2 except that instead of 4-(diphenylamino)benzaldehyde N-2,3-epoxypropyl-N-phenylhydrazone, 4-diethylaminobenzaldehyde N-2,3-epoxypropyl-N-phenyl hydrazone (3.60 g, 11.13 mmol) was used. The yield was 5.2 g (67.5 %), and the product had a melting point of 140.5-142 °C when crystallized from 10:1 v/v of 2-propanol and ether). A ¹HNMR spectrum (250 MHz, CDCl₃) yielded peaks at, ppm: 7.87 (d, *J*=8.8 Hz; 1H); 7.55 (s, 1H); 7.49 (s, 1H); 7.45 (d, *J*=5.7 Hz; 2H); 7.40-6.89 (m, 15H); 6.63 (d, *J*=8.9 Hz; 2H); 6.38 (d, *J*=8.8 Hz; 1H); 6.02 (s, 1H); 4.17 (m, 1H); 3.99-3.62 (m, 4H); 3.45 (m, 8H); 2.83 (d, *J*=6.5 Hz; 1H); 1.12 (m, 12H). An elemental analysis yielded, %: C 75.45; H 7.11; N 12.51. C₄₃H₅₀N₆O₂ has a calculated elemental distribution of, %: C 75.63; H 7.38; N 12.31.

Example 3 - Charge Mobility Measurements

[0092] This example describes the measurement of charge mobility for samples formed with the five charge transport compounds described in Example 2. Some samples included a polymer binder while other samples did not.

Sample 1

[0093] A mixture of 0.1 g of Compound 2 and 0.1 g of polyvinylbutyral (Aldrich 41,843-9) was dissolved in 2 ml of THF. The solution was coated on a polyester film with conductive aluminum layer by the dip roller method. After drying for 15 min. at 80°C temperature, a clear 10 μm thick layer was formed.

Sample 2

[0094] A 0.24 g quantity of Compound 3 was dissolved in 1 ml of THF. The solution was coated on the polyester film with conductive aluminum layer by the dip roller method. After drying for 15 min. at 80°C temperature, a clear 5 μm thick layer of good quality of Compound 3 was formed.

Sample 3

[0095] The sample was prepared according to the procedure for Sample 1, except that Compound 3 was used in place of Compound 2.

Sample 4

[0096] A thin layer of Compound 4 was prepared according to the procedure for Sample 2 with Compound 4 replacing Compound 3.

Sample 5

[0097] The sample was prepared according to the procedure for Sample 1, except that Compound 4 was used in place of Compound 2.

Sample 6

[0098] The sample was prepared according to the procedure for Sample 1, except Compound 5 was used in place of Compound 2.

Sample 7

[0099] The sample was prepared according to the procedure for Sample 1, except Compound 6 was used in place of Compound 2.

Mobility Measurements

[0100] Each sample was corona charged positively up to a surface potential U , illuminated with 2 ns long nitrogen laser light pulse and the hole mobility μ was determined as described in Kalade et al., "Investigation of charge carrier transfer in electrophotographic layers of chalcogenide glasses," Proceeding IPCS 1994: The Physics and Chemistry of Imaging Systems, Rochester, NY, pp. 747-752, incorporated herein by reference. This hole mobility measurement was repeated changing the charging regime and charging the sample to different U values, which corresponded to different electric field strength inside the layer E . This dependence was approximated by the formula

$$\mu = \mu_0 e^{\alpha \sqrt{E}}$$

Here E is electric field strength, μ_0 is the zero field mobility and α is Pool-Frenkel parameter. The mobility characterizing parameters μ_0 and α values as well as the mobility value at the 6.4×10^5 V/cm field strength as determined from these measurements are given in Table 1.

Table 1

Sample	μ_0 (cm ² /V·s)	μ (cm ² /V·s) at $6.4 \cdot 10^5$ V/cm	α (cm/V) ^{1/2}
1	$2.1 \cdot 10^{-9}$	$7.7 \cdot 10^{-8}$	0.0045
2	$7.0 \cdot 10^{-8}$	$6.7 \cdot 10^{-6}$	0.0057
3	$1.1 \cdot 10^{-9}$	$1.5 \cdot 10^{-7}$	0.0061
4	$5.7 \cdot 10^{-7}$	$2.9 \cdot 10^{-5}$	0.0049
5	$1.7 \cdot 10^{-8}$	$1.5 \cdot 10^{-6}$	0.0056
6	$7.1 \cdot 10^{-9}$	$3.9 \cdot 10^{-7}$	0.0050
7	$1.6 \cdot 10^{-8}$	$1.3 \cdot 10^{-6}$	0.0055

Example 4 - Ionization Potential Measurements

[0101] This example describes the measurement of the ionization potential for the five charge transport compounds described in Example 2.

[0102] To perform the ionization potential measurements, a thin layer of charge transport compound about 0.5 μ m thickness was coated from a solution of 2 mg of charge transport compound in 0.2 ml of tetrahydrofuran on a 20 cm² substrate surface. The substrate was polyester film with an aluminum layer over a methylcellulose sublayer of about 0.4 μ m thickness.

[0103] Ionization potential was measured as described in Grigalevicius et al., "3,6-Di(N-diphenylamino)-9-phenyl-carbazole and its methyl-substituted derivative as novel hole-transporting amorphous molecular materials," Synthetic Metals 128 (2002), p. 127-131, incorporated herein by reference. In particular, each sample was illuminated with monochromatic light from the quartz monochromator with a deuterium lamp source. The power of the incident light beam was $2.5 \cdot 10^{-8}$ W. The negative voltage of -300 V was supplied to the sample substrate. The counter-electrode with the 4.5×15 mm² slit for illumination was placed at 8 mm distance from the sample surface. The counter-electrode was connected to the input of the BK2-16 type electrometer, working in the open input regime, for the photocurrent measurement. A 10^{-15} - 10^{-12} amp photocurrent was flowing in the circuit under illumination. The photocurrent, I , was strongly dependent on the incident light photon energy $h\nu$. The $I^{0.5}=f(h\nu)$ dependence was plotted. Usually the dependence of the square root of photocurrent on incident light quanta energy is well described by linear relationship near the threshold.

[see references "Ionization Potential of Organic Pigment Film by Atmospheric Photoelectron Emission Analysis," Electrophotography, 28, Nr. 4, p. 364 (1989) by E. Miyamoto, Y. Yamaguchi, and M. Yokoyama; and "Photoemission in Solids," Topics in Applied Physics, 26, 1-103 (1978) by M. Cordona and L. Ley, both of which are incorporated herein by reference]. The linear part of this dependence was extrapolated to the hv axis and Ip value was determined as the photon energy at the interception point. The ionization potential measurement has an error of ± 0.03 eV. The ionization potential values are given in Table 2.

Table 2 -

Ionization Potential	
Compound	Ip (eV)
2	5.27
3	5.22
4	5.20
5	5.28
6	5.19

[0104] As understood by those skilled in the art, additional substitution, variation among substituents, and alternative methods of synthesis and use may be practiced within the scope and intent of the present disclosure of the invention. The embodiments above are intended to be illustrative and not limiting. Additional embodiments are within the claims. Although the present invention has been described with reference to particular embodiments, workers skilled in the art will recognize that changes may be made in form and detail without departing from the spirit and scope of the invention.

[0105] Attention is directed to all papers and documents which are filed concurrently with or previous to this specification in connection with this application and which are open to public inspection with this specification, and the contents of all such papers and documents are incorporated herein by reference.

[0106] All of the features disclosed in this specification (including any accompanying claims, abstract and drawings), and/or all of the steps of any method or process so disclosed, may be combined in any combination, except combinations where at least some of such features and/or steps are mutually exclusive.

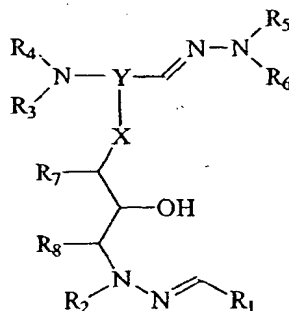
[0107] Each feature disclosed in this specification (including any accompanying claims, abstract and drawings) may be replaced by alternative features serving the same, equivalent or similar purpose, unless expressly stated otherwise. Thus, unless expressly stated otherwise, each feature disclosed is one example only of a generic series of equivalent or similar features.

[0108] The invention is not restricted to the details of the foregoing embodiment(s). The invention extends to any novel one, or any novel combination, of the features disclosed in this specification (including any accompanying claims, abstract and drawings), or to any novel one, or any novel combination, of the steps of any method or process so disclosed.

Claims

1. An organophotoreceptor comprising at least one photoconductive element comprising:

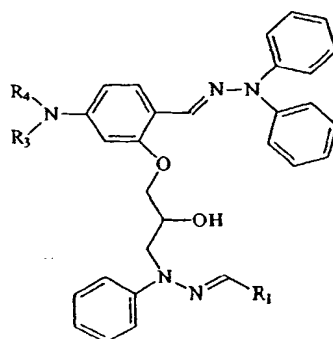
(a) a charge transport compound having the formula



where R_1 is a carbazole group, a julolidine group, or a p-(N,N-disubstituted)arylamine, R_2 , R_3 , R_4 , R_5 and R_6 are, independently, an alkyl group or an aryl group, R_7 and R_8 are, independently, hydrogen, an alkyl group, or an aryl group, X is oxygen, sulfur, or a NR' group where R' is hydrogen, an alkyl, or an aryl group, and Y is an aryl group; and
(b) a charge generating compound;

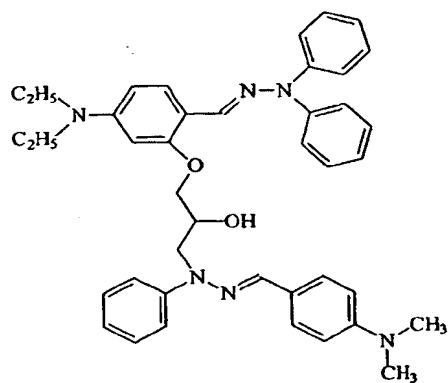
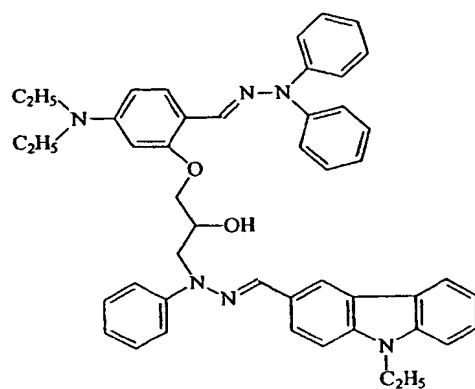
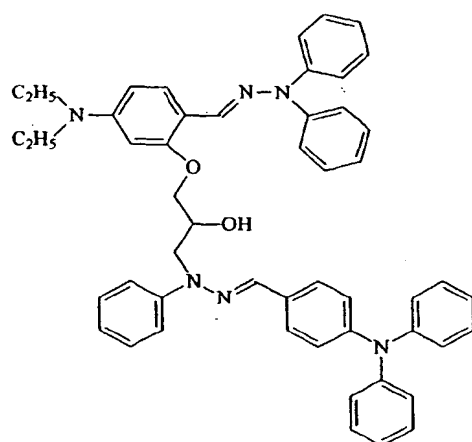
wherein the at least one photoconductive element is on an electrically conductive substrate.

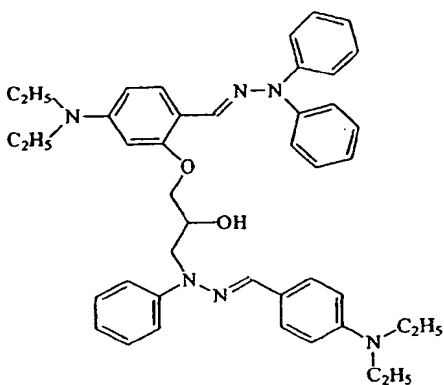
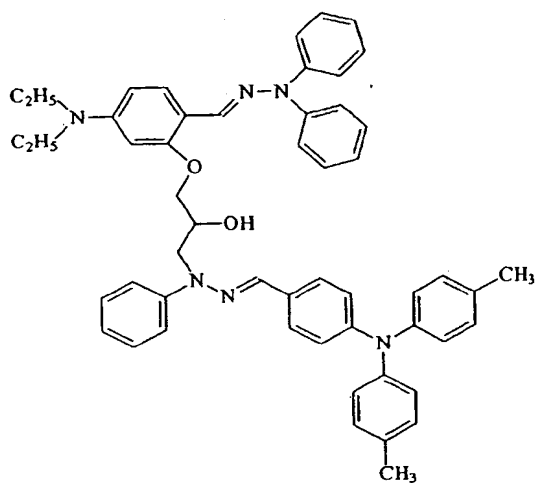
2. An organophotoreceptor according to claim 1 wherein the at least one photoconductive element further comprises an electron transport compound.
3. An organophotoreceptor according to either of claims 1 and 2 wherein the charge transport compound has the formula



where R_1 is a carbazole group, a julolidine group, or a p-(N,N-disubstituted)arylamine, and R_3 and R_4 are, independently, an alkyl group or an aryl group.

4. An organophotoreceptor according to any preceding claim wherein the at least one photoconductive element further comprises a binder.
5. An organophotoreceptor according to any preceding claim wherein the charge transport compound has a formula selected from the group consisting of the following:

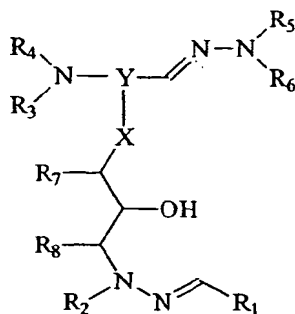




6. An electrophotographic imaging apparatus comprising:

- (a) a light imaging component; and
- (b) an organophotoreceptor oriented to receive light from the light imaging component, the organophotoreceptor comprising an electrically conductive substrate and at least one photoconductive element on the electrically conductive substrate, the photoconductive element comprising

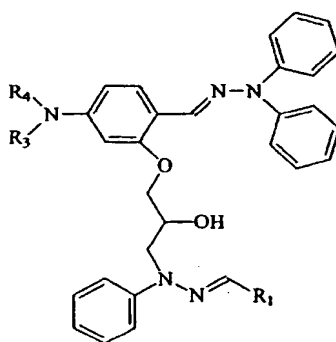
(i) a charge transport compound having the formula



where R_1 is a carbazole group, a julolidine group, or a p-(N,N-disubstituted)arylamine, R_2 , R_3 , R_4 , R_5 and R_6 are, independently, an alkyl group or an aryl group, R_7 and R_8 are, independently, hydrogen, an alkyl group, or an aryl group, X is oxygen, sulfur, or a NR' group where R' is hydrogen, an alkyl, or an aryl group, and Y is a aryl group; and

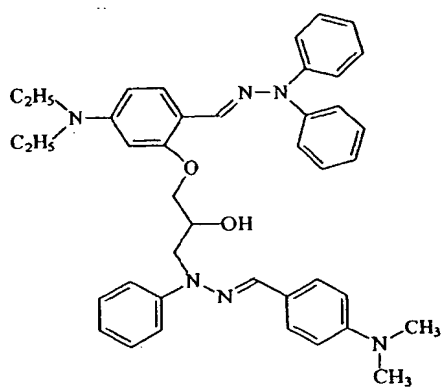
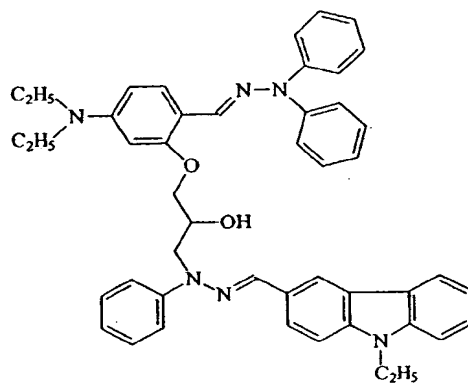
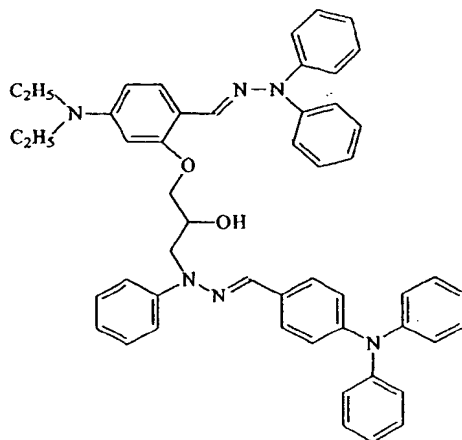
(ii) a charge generating compound.

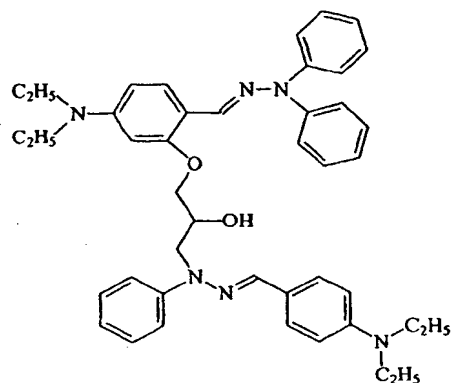
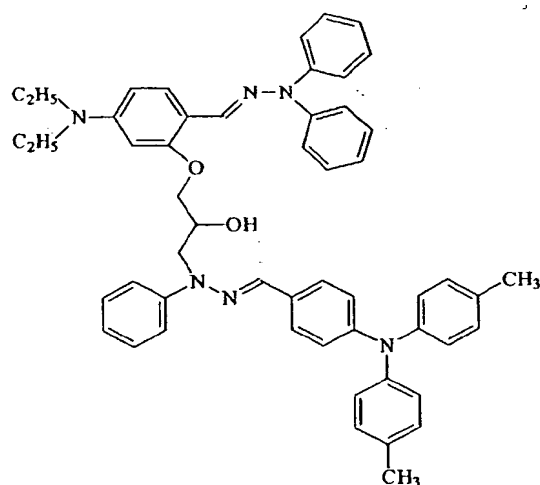
7. An electrophotographic imaging apparatus according to claim 6 wherein the charge transport compound has the formula



where R_1 is a carbazole group, a julolidine group, or a p-(N,N-disubstituted)arylamine, and R_3 and R_4 are, independently, an alkyl group or an aryl group.

8. An electrophotographic imaging apparatus according to either of claims 6 and 7 wherein the charge transport compound has a formula selected from the group consisting of the following:





9. An electrophotographic imaging apparatus according to any of claims 6 to 8 wherein the at least one photoconductive element further comprises an electron transport compound.

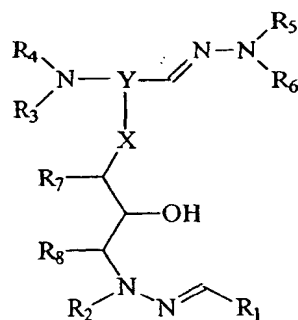
10. An electrophotographic imaging apparatus according to any of claims 6 to 9 wherein the at least one photoconductive element further comprises a binder.

11. An electrophotographic imaging apparatus according to any of claims 6 to 10 further comprising a liquid toner dispenser.

12. An electrophotographic imaging process comprising:

(a) applying an electrical charge to a surface of an organophotoreceptor comprising an electrically conductive substrate and at least one photoconductive element on the electrically conductive substrate, the photoconductive element comprising

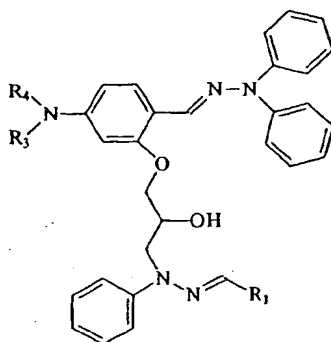
(i) a charge transport compound having the formula



where R_1 is a carbazole group, a julolidine group, or a p-(N,N-disubstituted)arylamine, R_2 , R_3 , R_4 , R_5 and R_6 are, independently, an alkyl group or an aryl group, R_7 and R_8 are, independently, hydrogen, an alkyl group, or an aryl group, X is oxygen, sulfur, or a NR' group where R' is hydrogen, an alkyl, or an aryl group, and Y is a aryl group; and
(ii) a charge generating compound;

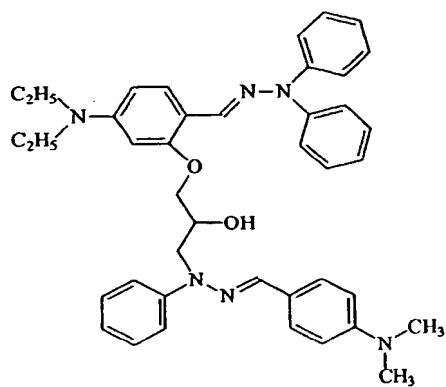
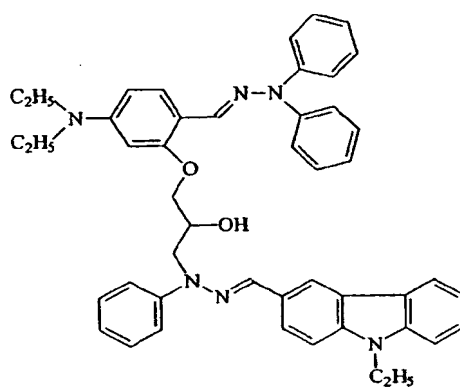
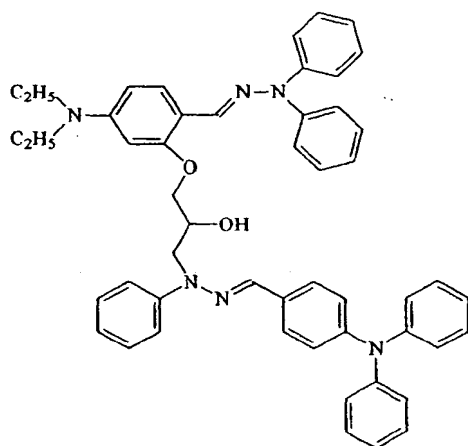
- (b) imagewise exposing the surface of the organophotoreceptor to radiation to dissipate charge in selected areas and thereby form a pattern of charged and uncharged areas on the surface;
- (c) contacting the surface with a toner to create a toned image; and
- (d) transferring the toned image to a substrate.

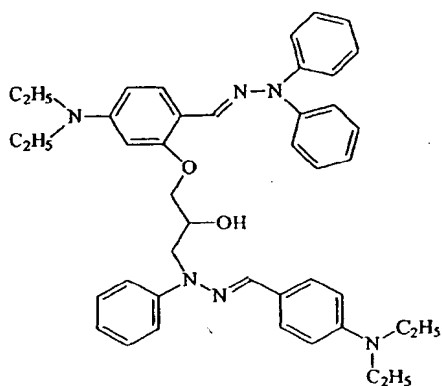
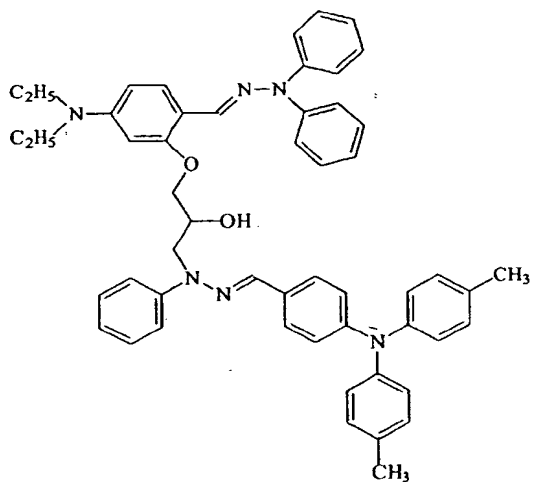
13. An electrophotographic imaging process according to claim 12 wherein the charge transport compound has the formula



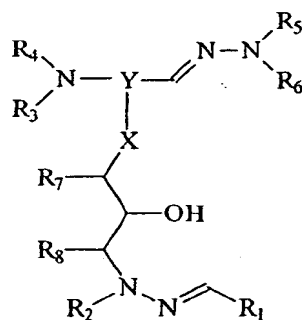
where R_1 is a carbazole group, a julolidine group, or a p-(N,N-disubstituted)arylamine, and R_3 and R_4 are, independently, an alkyl group or an aryl group.

14. An electrophotographic imaging process according to either of claims 12 and 13 wherein the charge transport compound has a formula selected from the group consisting of the following:



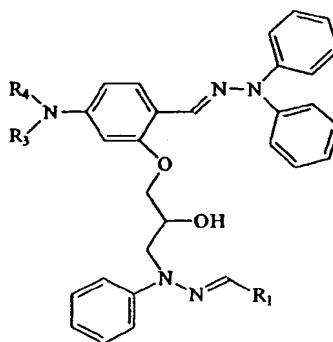


15. An electrophotographic imaging process according to any of claims 12 to 14 wherein the at least one photoconductive element further comprises an electron transport compound.
16. An electrophotographic imaging process according to any of claims 12 to 15 wherein the at least one photoconductive element further comprises a binder.
17. An electrophotographic imaging process according to any of claims 12 to 16 wherein the toner comprises a liquid toner comprising a dispersion of colorant particles in an organic liquid.
18. A charge transport compound having the formula



where R_1 is a carbazole group, a julolidine group, or a p-(N,N-disubstituted)arylamine, R_2 , R_3 , R_4 , R_5 and R_6 are, independently, an alkyl group or an aryl group, R_7 and R_8 are, independently, hydrogen, an alkyl group, or an aryl group, X is oxygen, sulfur, or a NR' group where R' is hydrogen, an alkyl, or an aryl group, and Y is a aryl group.

19. A charge transport compound according to claim 18 having the formula



where R_1 is a carbazole group, a julolidine group, or a p-(N,N-disubstituted)arylamine, and R_3 and R_4 are, independently, an alkyl group or an aryl group.

20. A charge transport compound according to either of claims 18 and 19 wherein the charge transport compound has a formula selected from the group consisting of the following:

